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NOTICE OF ALLOWANCE AND FEE(S) DUE

38199

7590

08/02/2010

HOWSON & HOWSON LLP / WYETH LLC 501 OFFICE CENTER DRIVE SUITE 210 FORT WASHINGTON, PA 19034 EXAMINER

BARHAM, BETHANY P

ART UNIT PAPER NUMBER

1615

DATE MAILED: 08/02/2010

A	APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
	10/574,210	08/29/2006	Sripriya Venkata Ramana Rao	AM-101457-1	5687

TITLE OF INVENTION: PANTOPRAZOLE MULTIPARTICULATE FORMULATIONS

APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	NO	\$1510	\$300	\$0	\$1810	11/02/2010

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. PROSECUTION ON THE MERITS IS CLOSED. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN THREE MONTHS FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. THIS STATUTORY PERIOD CANNOT BE EXTENDED. SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

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501 OFFICE CENTER DRIVE			ART UNIT	PAPER NUMBER
SUITE 210 FORT WASHING	TON, PA 19034		1615 DATE MAILED: 08/02/201	0

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment to date is 586 day(s). If the issue fee is paid on the date that is three months after the mailing date of this notice and the patent issues on the Tuesday before the date that is 28 weeks (six and a half months) after the mailing date of this notice, the Patent Term Adjustment will be 586 day(s).

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

	Application No.	Applicant(s)			
	10/574,210	VENKATA RAMANA RAO ET AL.			
Notice of Allowability	Examiner	Art Unit			
	BETHANY BARHAM	1615			
The MAILING DATE of this communication appearuments filed the Office or upon petition by the applicant. See 37 CFR 1.313 1. ☑ This communication is responsive to the amendments filed.	(OR REMAINS) CLOSED in to or other appropriate communing IGHTS. This application is suggested and MPEP 1308.	this application. If not included ication will be mailed in due course. THIS			
	. OH OO/ 1-1/ 10.				
2. ☑ The allowed claim(s) is/are <u>89-112</u> .					
 3. Acknowledgment is made of a claim for foreign priority ur a) All b) Some* c) None of the: 1. Certified copies of the priority documents have 2. Certified copies of the priority documents have 	e been received. e been received in Application	No			
 Copies of the certified copies of the priority do International Bureau (PCT Rule 17.2(a)). 	cuments have been received	in this national stage application from the			
* Certified copies not received:					
Applicant has THREE MONTHS FROM THE "MAILING DATE" noted below. Failure to timely comply will result in ABANDONN THIS THREE-MONTH PERIOD IS NOT EXTENDABLE. 4. A SUBSTITUTE OATH OR DECLARATION must be subm	IENT of this application. itted. Note the attached EXAN	MINER'S AMENDMENT or NOTICE OF			
INFORMAL PATENT APPLICATION (PTO-152) which give	. , -	declaration is deficient.			
5. CORRECTED DRAWINGS (as "replacement sheets") must be submitted.					
(a) ☐ including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached1) ☐ hereto or 2) ☐ to Paper No./Mail Date					
(b) ☐ including changes required by the attached Examiner's Paper No./Mail Date Identifying indicia such as the application number (see 37 CFR 1	s Amendment / Comment or i				
each sheet. Replacement sheet(s) should be labeled as such in t					
6. DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL.					
Attachment(s) 1. ☐ Notice of References Cited (PTO-892) 2. ☐ Notice of Draftperson's Patent Drawing Review (PTO-948) 3. ☑ Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date See Continuation Sheet 4. ☐ Examiner's Comment Regarding Requirement for Deposit of Biological Material	6. ☐ Interview Sur Paper No./M 7. ☑ Examiner's A 8. ☑ Examiner's S	ormal Patent Application mmary (PTO-413), lail Date mendment/Comment statement of Reasons for Allowance			
	9.				

Continuation of Attachment(s) 3. Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date: 01/08/10 and 03/21/07.

DETAILED ACTION

Status of the Claims

Receipt is acknowledged of the Applicant's Response and Terminal Disclaimers filed on 06/14/10. IDS's filed on 01/08/10 and 03/21/07 which were considered previously, now both documents 92 and 84 respectively have been reconsidered with the dates of the documents noted on the IDS. Document 92 dated 02/12/08 and document 84 dated 10/01. Claims 89-112 are pending.

Terminal Disclaimers

The terminal disclaimer filed on 06/14/10 disclaiming the terminal portion of any patent granted on this application which would extend beyond the expiration date of US Patents 7,544,370; 7,553,498 and 7,550,153 have been reviewed and are accepted. The terminal disclaimers have been recorded.

Examiner's Amendment

Authorization for this examiner's amendment was given in a telephone interview with Cathy Kodroff on 07/06/10. Claims 89, 100, 101 and 106 have been amended as follows:

Claim 89: Pantoprazole multiparticulates having reduced release under gastric conditions and fast release at neutral pH, wherein each of said multiparticulates comprises: a spheroid core [consisting of] comprising about 20 % w/w to about 45%

w/w of a pantoprazole salt or a hydrate thereof, as the sole active component in the multiparticulate and [one or more excipients comprising] about 25% to about 30% w/w microcrystalline cellulose, about 4% to about 6% w/w polysorbate 80, about 14% to about 16% w/w crospovidone, about 0.5 to about 2% w/w hydroxypropyl methylcellulose, about 5% to about 8% w/w sodium carbonate, and about 1 to about 2% w/w water;

an initial seal coat comprising hydroxypropylmethyl cellulose on the spheroid core; and

an enteric coat on the initial seal coat,

wherein said multiparticulates have an average diameter of about 0.7 mm to about 1.25 mm.

Claim 100: Pantoprazole multiparticulates having reduced release under gastric conditions and fast release at neutral pH, wherein each of said multiparticulates comprises: a spheroid core [consisting of] <u>comprising</u> a pantoprazole salt or a hydrate thereof in an amount of about 40% w/w free pantoprazole, as the sole active component in the multiparticulate and [one or more excipients comprising] about 25% to about 30% w/w microcrystalline cellulose, about 4% to about 6% w/w polysorbate 80, about 14% to about 16% w/w crospovidone, about 0.5 to about 2% w/w hydroxypropyl methylcellulose, about 5% to about 8% w/w sodium carbonate, and about 1 to about 2% w/w water; an initial seal coat comprising hydroxypropylmethyl cellulose on the

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spheroid core; and an enteric coat on the initial seal coat, wherein said multiparticulates have an average diameter of about 0.7 mm to about 1.25 mm.

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Claim 101: A product comprising a plurality of pantoprazole multiparticulates, each comprising a spheroid core [consisting of] <u>comprising</u> a pantoprazole salt or a hydrate thereof in an amount of about 40% w/w free pantoprazole, <u>as the sole active component in the multiparticulate</u> and [one or more excipients comprising] about 25% to about 30% w/w microcrystalline cellulose, about 4% to about 6% w/w polysorbate 80, about 14% to about 16% w/w crospovidone, about 0.5 to about 2% w/w hydroxypropyl methylcellulose, about 5% to about 8% w/w sodium carbonate, and about 1 to about 2% w/w water; an initial seal coat comprising hydroxypropylmethyl cellulose on the spheroid core; and an enteric coat on the initial seat coat, wherein said multiparticulates have an average diameter of about 0.7 mm to about 1.25 mm.

Claim 106: A method of producing a multiparticulate formulation of pantoprazole having reduced release under gastric conditions and fast release at neutral pH, said method comprising: producing a spheroid core [consisting of] comprising a pantoprazole salt or a hydrate thereof in an amount of about 40% w/w free, as the sole active component in the multiparticulate and [one or more excipients comprising] about 25% to about 30% w/w microcrystalline cellulose, about 4% to about 6% w/w polysorbate 80, about 14% to about 16% w/w crospovidone, about 0.5 to about 2% w/w hydroxypropyl methylcellulose, about 5% to about 8% w/w sodium carbonate, and about 1 to about 2

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% w/w water, via extrusion and spheronization; applying an initial seal coat comprising hydroxypropyl methyl cellulose to the spheroid core; applying an enteric coating to the initial seal coated spheroid core, said enteric coating comprising a copolymer of methacrylic acid and methacrylates; and optionally applying a final seal coat to the enteric-coated spheroid core, said final seal coat being about 1 wt% of the multiparticulate; wherein the multiparticulates have an average diameter of about 0.7 mm to about 1.25 mm.

Reasons for allowance

The following is an examiner's statement of reasons for allowance:

The prior art of record does not teach or suggest a spheroid core with pantoprazole salt or a hydrate thereof as the sole active component with all the specific excipients and specific ranges as instant claimed.

Therefore the claims are allowable over the prior art of record.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

Conclusion

Claims 89-112 are allowed.

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Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Bethany P. Barham whose telephone number is 571-272-6175. The examiner can normally be reached on M-F from 8:30am to 5pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Robert A. Wax, can be reached on 571-272-0623. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Bethany Barham Examiner, Art Unit 1615 /Robert A. Wax/ Supervisory Patent Examiner, Art Unit 1615